



# STIC Search Report

## Biotech-Chem Library

STIC Database Tracking Number: 1635402

**TO:** Janet Epps-Ford  
**Location:** REM-2C05/2C18  
**Art Unit:** 1635  
**Thursday, December 02, 2004**  
**Case Serial Number:** 09/551494

**From:** Paul Schulwitz  
**Location:** Biotech-Chem Library  
**REM-1A65**  
**Phone:** (571)272-2527  
  
**[paul.schulwitz@uspto.gov](mailto:paul.schulwitz@uspto.gov)**

### Search Notes

Examiner Epps-Ford,

See attached results.

If you have any questions about this search feel free to contact me at any time.

Thank you for using STIC search services!

Paul Schulwitz  
Technical Information Specialist  
STIC Biotech/Chem Library  
(571)272-2527

**Schulwitz, Paul**

**From:** Epps-Ford, Janet  
**Sent:** Wednesday, December 01, 2004 12:33 PM  
**To:** Schulwitz, Paul  
**Subject:** Sequence and Word-Search

Hi there, I was wondering if I can get the following claim language searched:

Application 09/551,494

An RNA viral vector comprising the nucleotide sequence of SEQ ID NO: 5 from the nucleotide at position 5430 to the nucleotide at position 5505.

*Thanks,  
Janet L. Epps-Ford, Ph.D.  
Art Unit 1635  
Mailbox: Remsen 2C18  
Office: Remsen 2C05  
Phone: 571-272-0757  
Fax: 571-273-0757*

GenCore version 5.1.6  
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## Om nucleic - nucleic search, using sw model

Run on: December 1, 2004, 20:12:59 ; Search time 408 Seconds  
(without alignments)

977.833 Million cell updates/sec

Title: US-09-551-494-5\_COPY\_5430\_5505

Perfect score: 76  
Sequence: 1 gtgacagacggctcgccat.....tgaagtaccaatggctgtga 76

Scoring table: IDENTITY\_NUC  
Gappen 10<sup>-3</sup>, Gapext 1.0

Searched: 4134886 seqs, 2624710521 residues

Total number of hits satisfying chosen parameters: 8269772

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match 0%  
Maximum Match 100%

Listing first 45 summaries

Database :

N\_Geneseq\_23Sep04: \*  
1: geneseqn1980s: \*  
2: geneseqn1990s: \*  
3: geneseqn2000s: \*  
4: geneseqn2010s: \*  
5: geneseqn2001s: \*  
6: geneseqn2002s: \*  
7: geneseqn2002s8: \*  
8: geneseqn2003s: \*  
9: geneseqn2003s8: \*  
10: geneseqn2004s: \*  
11: geneseqn2003s8: \*  
12: geneseqn2004s: \*

Prob. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

## SUMMARIES

Result No. Score Query

Match Length DB ID

Description

RESULT 1  
AAC62372

ID AAC62372 standard; DNA; 6355 BP.

XX AAC62372;

XX DT 19-MAR-2001 (first entry)

XX CDNA sequence of the genome of tobacco mosaic virus-U2.

XX Plant phenotype; gene trait; Nicotiana; Oryza sativa; Zea mays; Brassica;

XX Gossypium; Triticum; Arabidopsis; Petunia; herbicide; transgenic plant;

XX tobacco mosaic virus; TMV; tobacco mosaic virus; TMV; helper virus; ss-

XX OG Tobacco mosaic virus.

XX PN WO200063397-A2.

XX PD 26-OCT-2000.

XX XY 17-APR-2000; 2000WO-EP003521.

XX PR 20-APR-1999; 99US-00234022.

XX (AVET ) AVENTIS CROPSCIENCE NV.

XX Meulewaeter F, Cornelissen M, Jacobs J, Van Eldik G, Metzlaff M;

XX WPI: 2000-687182-67.

XX Identifying and isolating genes involved in determining the trait or

XX phenotype of plant species, by infecting plants with gene silencing

XX constructs targeted to the gene, and identifying plants with altered

XX traits.

Example 1; Page 53-56; 64pp; English.

XX The specification describes a method for isolating genes that determine a trait or phenotype of a plant species. The method comprises identifying a set of nucleic acids of genes correlated with the trait, creating a library of gene silencing constructs in a viral RNA vector, targeting the gene silencing constructs to the nucleic acid set, infecting a collection of individual plants with these, identifying plants with altered traits or phenotype, and isolating genes of the invention. The method is useful.

CC for isolating genes involved in the determination of trait or a phenotype of a plant such as *Nicotiana*, *Oryza sativa*, *Zea mays*, *Brassica*, *Gossypium*, *Triticum*, *Arabidopsis* or *Petunia*. The method is also useful for modulating the expression of selected nucleic acid sequences and for validating the function of a nucleic acid sequence whose expression is correlated with the presence or absence of a specific trait in plants, but with otherwise unknown function. The method is also useful for developing agriculturally useful products such as herbicides or transgenic plants. The present sequence represents the cDNA sequence of the genome of tobacco mosaic virus (TMV)-U2. The sequence was used to construct a plasmid vector for the synthesis of an infective hybrid tobacco mosaic virus (TMV) satellite tobacco necrosis virus (STNV) helper virus RNA. This helper virus is used in the method of the invention.

XX Sequence 6355 BP; 1933 A; 1112 C; 1489 G; 1821 T; 0 J; 0 Other; SQ Best Local Similarity 100.0%; Pred. No. 2.2e-16; Length 6355; Matches 76; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTGACAGAGGGCTGGCCATTGACTGAAAGGTGTTAGGAGTCGCGATGAA 60  
ID AD126338 standard; DNA; 769 52.  
XX  
AC AD126338;  
XX  
DB 5430 GTGACAGAGGGCTGGCCATTGACTGAAAGGTGTTAGGAGTCGCGATGAA 5489  
QY 61 GTTACCAATGGCTGTGA 76  
ID 5490 GTTACCAATGGCTGTGA 5505  
XX  
AC  
DB

RESULT 2

AC085005  
ID ACC85005 standard; DNA; 6355 BP.  
XX  
AC ACC85005;  
XX  
DT 13-OCT-2003 (first entry)  
XX  
DE TMV-U2 genome nucleotide sequence.  
XX  
KW Inhibitory RNA; viral RNA vector; coat protein; TMV; U2; gene; ds.  
XX  
OS Tobacco mosaic virus.  
XX  
PN WO2003052108-A2.  
XX  
PD 26-JUN-2003.  
XX  
PP 05-DEC-2002; 2002WO-EP013964.  
XX  
PR 18-DEC-2001; 2001US-0340498P.  
XX  
PA (PARB ) BAYER BIOSCIENCE NV.  
XX  
PT Metzlaff MH, Gosselle VML, Meulewaeter F, Fache ICA,  
DR WPI; 2003-523529/49.  
XX  
PT Introducing inhibitory RNA into a plant cell comprises providing a viral RNA vector derived from a satellite RNA virus that encodes a coat protein, and infecting a plant with the viral RNA vector and a corresponding helper virus.

XX  
PS Example; Page 79-82; 86pp; English.

CC The invention relates to introducing inhibitory RNA into a plant cell. The method involves providing a viral RNA vector derived from a satellite RNA virus having a sequence that encodes a coat protein, and infecting a plant with the viral RNA vector and a corresponding helper virus. The methods and viral RNA vectors are useful in introducing inhibitory RNA into plant cells. These may be used to determine or validate the function of isolated nucleic acid sequences in plants. The present sequence represents the nucleotide sequence of the genome of tobacco mosaic virus (TMV)-U2.

XX Sequence 6355 BP; 1933 A; 1112 C; 1489 G; 1821 T; 0 J; 0 Other; SQ Best Local Similarity 100.0%; Pred. No. 2.2e-16; Length 6355; Matches 76; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GTGACAGAGGGCTGGCCATTGACTGAAAGGTGTTAGGAGTCGCGATGAA 60  
ID 5430 GTGACAGAGGGCTGGCCATTGACTGAAAGGTGTTAGGAGTCGCGATGAA 5489  
QY 61 GTTACCAATGGCTGTGA 76  
ID 5490 GTTACCAATGGCTGTGA 5505  
XX  
AC  
DB

RESULT 3

AD126338  
ID AD126338 standard; DNA; 769 52.  
XX  
AC AD126338;  
XX  
DT 22-APR-2004 (first entry)  
XX  
DE Novel endonuclease Res I-related clone DNA 3.  
XX  
KW endonuclease; molecular biology; Plant propagation; phenotypic trait; herbicide tolerance; heat tolerance; cold tolerance; drought; salinity; osmotic stress; pest resistance; insect; nematode; arachnid; fungal; bacterial; viral; enzyme production; secondary metabolite; male sterility; female sterility; dwarfness; early maturity; Res I; ds.  
XX  
OS Tobacco mosaic virus.  
XX  
PN US2003148315-A1.  
XX  
PD 07-AUG-2003.  
XX  
PF 01-AUG-2002; 2002US-00211079.  
XX  
ER 01-FEB-2002; 2002US-0353722P.  
PR 24-MAR-2002; 2002US-00098155.  
XX  
PA (PADG/ ) PADGETT H S.  
XX  
PA (VAEW/ ) VAEHONGS A A.  
XX  
PI Padgett HS, Vaehonges AA;  
XX  
DR WPI; 2003-897548/82.  
XX  
PT New nucleic acid molecule encoding endonucleases, useful in molecular biology, specifically to generating populations of related nucleic acid molecules, and in plant propagation with useful phenotypic traits.  
XX  
PS Example 15; Fig 8; 46pp; English.

CC This invention relates to a novel endonuclease (Res I) nucleic acid molecule which comprises a fully defined sequence of 899 bp given in the specification. The methods and compositions of the present invention are useful in molecular biology, and more specifically to generating populations of related nucleic acid molecules. They may also be used in plant propagation with useful phenotypic traits, such as improved tolerance to herbicides, improved tolerance to extremes of heat or cold, insects, nematodes or arachnids or diseases (fungal, bacterial or viral), production of enzymes or secondary metabolites, male or female sterility, dwarfness and early maturity. The present sequence is that of a clone which was derived during the exemplification of the invention.

XX Sequence 769 BP; 247 A; 102 C; 201 G; 219 T; 0 U; 0 Other; SQ Best Local Similarity 94.7%; Pred. No. 2.3e-14; Length 769;



Query Match 91.6%; Score 69; DB 12; Length 769;  
 Best Local Similarity 94.7%; Pred. No. 2; 3e-14; Indels 0; Gaps 0;  
 Matches 72; Conservative 0; Mismatches 4;

QY 1 GTGACAGACGGCTGCCAATTGAACTCTGAAAGGTGTAGGAGTCCTGGATGAA 60  
 DB 541 GTACAGACGGCTGCCAATTGAACTCTGAAAGGTGTAGGAGTCCTGGATGAA 600  
 QY 61 GTACCAATGGCTGTA 76  
 DB 601 GTACCAATGGCTGTA 616

RESULT 6  
 AD126344/C  
 ID AD126344 standard; DNA; 772 BP.  
 XX  
 AC AD126344;  
 XX  
 DT 22-APR-2004 (first entry)  
 XX  
 DE Novel endonuclease Res I-related clone DNA 9.  
 XX  
 KW endonuclease; molecular biology; plant propagation; phenotypic trait;  
 KW herbicide tolerance; heat tolerance; cold tolerance; drought; salinity;  
 KW osmotic stress; pest resistance; insect; nematode; arachnid; fungal;  
 KW bacterial; viral; enzyme production; secondary metabolite;  
 KW male sterility; female sterility; dwarfness; early maturity; Res I; ds.  
 XX  
 OS Tomato mosaic virus.  
 XX  
 US2003148315-A1.  
 XX  
 PD 07-AUG-2003.  
 XX  
 PP 01-AUG-2002; 2002US-00211079.  
 XX  
 PR 31-FEB-2002; 2002US-0353722P.  
 PR 14-MAR-2002; 2002US-00098155.  
 XX  
 PA (PADG/) PADGETT H S.  
 PA (VAEW/) VAEWHONGS A A.  
 PI Padgett HS, Vaewhongs AA;  
 XX  
 WPI; 2003-897548/82.

XX  
 PT New nucleic acid molecule encoding endonucleases, useful in molecular  
 PT biology, specifically to generating populations of related nucleic acid  
 PT molecules, and in plant propagation with useful phenotypic traits.  
 XX  
 PS Example 15; Fig 14; 46pp; English.

XX  
 CC This invention relates to a novel endonuclease (Res I) nucleic acid  
 CC molecule which comprises a fully defined sequence of 899 bp given in the  
 CC specification. The methods and compositions of the present invention are  
 CC useful in molecular biology, and more specifically to generating  
 CC populations of related nucleic acid molecules. They may also be used in  
 CC plant propagation with useful phenotypic traits, such as improved  
 CC tolerance to herbicides, improved tolerance to extremes of heat or cold,  
 CC drought, salinity or osmotic stress, improved resistance to pests  
 CC (insects, nematodes or arachnids) or diseases (fungal, bacterial or  
 CC viral); production of enzymes or secondary metabolites, male or female  
 CC sterility; dwarfness and early maturity. The present sequence is that of  
 CC a clone which was derived during the exemplification of the invention.  
 XX  
 SQ Sequence 772 BP; 227 A; 196 C; 109 G; 240 T; 0 U; 0 Other;

Query Match 89.5%; Score 68; DB 10; Length 772;  
 Best Local Similarity 93.4%; Pred. No. 8.2e-14; Indels 3; Gaps 0;  
 Matches 71; Conservative 3; Mismatches 5;

QY 1 GTGACAGACGGCTGCCAATTGAACTCTGAAAGGTGTAGGAGTCCTGGATGAA 60  
 DB 232 GTACAGACGGCTGCCAATTGAACTCTGAAAGGTGTAGGAGTCCTGGATGAA 173  
 QY 61 GTACCAATGGCTGTA 76  
 DB 172 GTACCAATGGCTGTA 157

RESULT 7  
 ADM68457/C  
 ID ADM68457 standard; DNA; 772 BP.  
 XX  
 AC ADM68457;  
 XX  
 DT 03-JUN-2004 (first entry)  
 XX  
 DE Mosaic virus movement protein gene GRAMMR clone #11.  
 XX  
 KW ds; mismatch endonuclease; endonuclease; gene shuffling technology;  
 KW single nucleotide polymorphism; cancer susceptibility; gene;  
 KW sequence variation redistribution; movement protein; gene.  
 XX  
 OS Tomato mosaic virus.  
 XX  
 DN US2003157682-A1.  
 XX  
 PD 21-AUG-2003.  
 XX  
 PP 31-JAN-2003; 2003US-00356708.  
 XX  
 PR 01-FEB-2002; 2002US-0353722P.  
 PR 14-MAR-2002; 2002US-00098155.  
 PR 01-AUG-2002; 2002US-00211079.  
 XX  
 PA (PADG/) PADGETT H S.  
 PA (VAEW/) VAEWHONGS A A.  
 PA (VOJD/) VOJDANI F S.  
 PA (SMIT/) SMITH M L.  
 PA (LIND/) LINDBO J A.  
 PA (FITZ/) FITZMAURICE W P.  
 XX  
 PA Padgett HS, Vaewhongs AA, Vojdani FS, Smith ML, Jindbo JA;  
 PI Fitzmaurice WP;  
 XX  
 DR WPI; 2003-766176/72.

XX  
 PT Making a mismatch endonuclease, useful in gene shuffling and in detection  
 PT of single nucleotide polymorphisms, comprises transfecting a host with a  
 PT recombinant viral vector including a polynucleotide encoding a mismatch  
 PT endonuclease.  
 XX  
 PS Example 14; SEQ ID NO 26; 79pp; English.

XX  
 CC The invention relates to a method of making a mismatch endonuclease  
 CC enzyme comprising transfecting a host plant, animal, yeast, fungus or  
 CC bacterium with a recombinant viral vector that encodes a polynucleotide  
 CC sequence for a mismatch endonuclease, growing the host so that the  
 CC polynucleotide is expressed, and extracting the mismatch endonuclease  
 CC enzyme from the host. The method is useful for making mismatch  
 CC endonuclease enzymes, for obtaining peptides and polynucleotides with  
 CC desired functional properties and for detecting mutations. The mismatch  
 CC endonuclease enzymes are useful in gene shuffling technology for  
 CC developing new genes, in detecting single nucleotide polymorphisms for  
 CC e.g. detecting evidence of cancer susceptibility, or in redistributing  
 CC sequence variations between non-identical polynucleotide sequences. The  
 CC present sequence represents a mosaic virus movement protein gene GRAMMR  
 CC clone.

XX  
 Sequence 772 BP; 227 A; 196 C; 109 G; 240 T; 0 U; 0 Other;  
 CC  
 CC

Query Match 89.5%; Score 68; DB 11; Length 772;

Best local similarity 93.4%; Pred. No. 8.2e-14; Matches 71; Conservative 0; Mismatches 5; Indels 0; Gaps 0; Query Match 89.5%; Score 68; DB 12; Length 772;

QY	1	GTGACAGAGCTGCTGCCAATGAACTGAAAGGTGTTGAGGAGTGTGATGAA	60
Db	232	GTAAAGAGACGCTGCCAATGAACTGAAAGGTGTTGAGGAGTGTGATGAA	173
QY	61	GTACCAATGCTGTGA	76
Db	172	GTACCAATGCTGTGA	157

RESULT 8

ADP26610/c	ID	ADP26610	standard; DNA; 772 BP.
XX	AC	ADP26610;	
XX	AC	ADP26610;	
XX	DT	26-AUG-2004	(first entry)
DE	DE	Heteroduplex DNA #15.	
KW	KW	Sequence variation; heteroduplex; transcription; DNA integration;	
OS	KW	ribozyme expression; gene; ds.	
XX	OS	Synthetic.	
XX	PN	US2004110130-A1.	
XX	PD	10-JUN-2004.	
XX	PP	25-OCT-2002; 2002US-00280913.	
XX	PR	02-FEB-2001; 2001US-0266386P.	
PR	PR	14-FEB-2001; 2001US-0268785P.	
PR	PR	01-FEB-2002; 2002US-0006690.	
PR	PR	08-AUG-2002; 2002US-0402342P.	
XX	PA	(LARG->) LARGE SCALE BIOLOGY CORP.	
PT	Padgett HS, Lirabo JA, Fitzmaurice WP;		
XX	DR	WPI; 2004-440326/41.	
PT	Redistributing sequence variations between non-identical polynucleotide sequences, useful for generating improved polynucleotide having a desired characteristic, comprises making a heteroduplex and introducing a nick.		
PT	Example 15; SEQ ID NO 26; 75pp; English.		
XX	CC	The invention relates to an in vitro method of redistributing sequence variations between non-identical polynucleotide sequences, comprising making a heteroduplex polynucleotide from two non-identical, complementary polynucleotides, introducing a nick in the second strand at or near a base pair mismatch site, removing the mismatched base(s) from the mismatch site where the nick occurred and using the first strand as a template to replace the removed base(s) with bases that complement the base(s) in the first strand. The invention also relates to an in vitro method of making a population of sequence variants from a heteroduplex polynucleotide sequence, obtaining a polynucleotide sequence encoding a desired functional property and identifying a reassembled DNA molecule encoding a protein with a desired functional property. The method is useful for generating an improved polynucleotide sequence or a population of improved polynucleotide sequences possessing at least one desired phenotypic characteristic (e.g., promotes transcription of linked polynucleotides), where such polynucleotides are useful for expression from a plant, animal, fungal, yeast, or bacterial expression vector, for integration to form a transgenic plant, animal or microorganism, and for expression of a ribozyme. This sequence represents DNA used in the scope of the invention.	
XX	SQ	Sequence 772 BP; 227 A; 196 C; 109 G; 240 T; 0 U; 0 Other;	

Query Match 89.5%; Score 68; DB 12; Length 772;

QY	1	GTGACAGAGCTGCTGCCAATGAACTGAAAGGTGTTGAGGAGTGTGATGAA	60
Db	232	GTAAAGAGACGCTGCCAATGAACTGAAAGGTGTTGAGGAGTGTGATGAA	173
QY	61	GTACCAATGCTGTGA	76
Db	172	GTACCAATGCTGTGA	157

RESULT 9

AAC62379	ID	AAC62379	standard; DNA; 411 BP.
XX	AC	AAC62379;	
XX	AC	AAC62379;	
XX	DT	19-MAR-2001	(first entry)
DE	Origin of assembly (CAS) of a tobacco mosaic virus (TMV) -U2.		
KW	KW	Plant phenotype; gene trait; Nicotiana; Cryza sativa; Zea mays; Brassica; Gossypium; Triticum; Arabidopsis; Petunia; herbicide; transgenic plant; tobacco necrosis virus; TMV; tobacco mosaic virus; TMV; helper virus; origin of assembly; ss.	
XX	OS	Tobacco mosaic virus.	
XX	PN	W020063397-A2.	
XX	PD	26-OCT-2000.	
XX	PF	17-APR-2000; 2000WO-EP003521.	
XX	PR	20-APR-1999; 99US-00294022.	
XX	PA	(AVET ) AVENTIS CROPSCIENCE NV.	
XX	PI	Medlewaeter F, Cornelissen M, Jacobs J, van Eldik G, Metzlaaff M;	
XX	DR	WPI; 2000-687182/67.	
PT	Identifying and isolating genes involved in determining the trait or phenotype of plant species, by infecting plants with gene silencing constructs targeted to the gene, and identifying plants with altered traits.		
PT	Example 1; Page 63; 64pp; English.		

Query Match 89.5%; Score 68; DB 12; Length 772;

QY	1	GTGACAGAGCTGCTGCCAATGAACTGAAAGGTGTTGAGGAGTGTGATGAA	60
Db	232	GTAAAGAGACGCTGCCAATGAACTGAAAGGTGTTGAGGAGTGTGATGAA	173
QY	61	GTACCAATGCTGTGA	76
Db	172	GTACCAATGCTGTGA	157

RESULT 9

AAC62379	ID	AAC62379	standard; DNA; 411 BP.
XX	AC	AAC62379;	
XX	AC	AAC62379;	
XX	DT	19-MAR-2001	(first entry)
DE	Origin of assembly (CAS) of a tobacco mosaic virus (TMV) -U2.		
KW	KW	Plant phenotype; gene trait; Nicotiana; Cryza sativa; Zea mays; Brassica; Gossypium; Triticum; Arabidopsis; Petunia; herbicide; transgenic plant; tobacco necrosis virus; TMV; tobacco mosaic virus; TMV; helper virus; origin of assembly; ss.	
XX	OS	Tobacco mosaic virus.	
XX	PN	W020063397-A2.	
XX	PD	26-OCT-2000.	
XX	PF	17-APR-2000; 2000WO-EP003521.	
XX	PR	20-APR-1999; 99US-00294022.	
XX	PA	(AVET ) AVENTIS CROPSCIENCE NV.	
XX	PI	Medlewaeter F, Cornelissen M, Jacobs J, van Eldik G, Metzlaaff M;	
XX	DR	WPI; 2000-687182/67.	
PT	Identifying and isolating genes involved in determining the trait or phenotype of plant species, by infecting plants with gene silencing constructs targeted to the gene, and identifying plants with altered traits.		
PT	Example 1; Page 63; 64pp; English.		

The specification describes a method for isolating genes that determine a trait or phenotype of a plant species. The method comprises identifying a set of nucleic acids of genes correlated with the trait; creating a library of gene silencing constructs to the nucleic acid set; infecting the gene silencing constructs to the nucleic acid set; infecting a collection of individual plant with these, identifying plants with altered traits or phenotype, and isolating genes of the invention. The method is useful for isolating genes involved in the determination of trait or a phenotype of a plant such as *Nicotiana*, *Oryza sativa*, *Zea mays*, *Brassica*, *Gossypium*, *Triticum*, *Arabidopsis* or *Petunia*. The method is also useful for modulating the expression of selected nucleic acid sequences and for validating the function of a nucleic acid sequence whose expression is correlated with the presence or absence of a specific trait in plants, but with otherwise unknown function. The method is also useful for developing agronomically useful products such herbicides or transgenic plants. The present sequence is an origin of assembly (OAs) of a tobacco mosaic virus (TMV)-U2. The sequence is used to construct infective hybrid tobacco mosaic virus (TMV)/tobacco necrosis virus (TNV) vectors, for use in the method of the invention

Sequence 411 BP; 140 A; 70 C; 93 G; 108 T; 0 U; 0 Other;

us-09-551-494-5 copy 5430 5505.rng

Query Match	77.1%	Score 58.6;	DB 3;	Length 411;
Best Local Similarity	93.8%	Pred. 1.4e-10;		
Matches	61;	Conservative 0;	Mismatches 4;	Indels 0;
2y				
12	CTGGCCATTGAACTGACTGAGGTTGAGGAGTCGTTGAGGATGAACTTGGC	71		
3b	CTGGCCATTGAACTGACTGAGGTTGAGGAGTCGTTGAGGATGAACTTGGC	62		

QY	1	GTGACAGAGCGGCTGCGCAATTGAACTCACTGAAAGGTTT3AGGAACTGTGAGTGA	60
Db	229	GTGAACTGGACGACCCATGGAACCTTCAAGAAGAGTGTGATGAGTCATAGTGA	170
QY	61	GTACCAATGGCTGTA	76
Db	159	GTACCAATGGCTGTA	154

Db	63	TGTGA	67
		/2 1G1GA 1    1	/6

RESULT 11  
ADM68456/c  
ID ADM68456 standard: DNA: 769 BP.

HW42034411  
ADT226343 standard; DNA; 769 BP.  
XX  
AC  
ADT26343;  
XX  
DT  
XX  
22-APR-2004 (first entry)  
Novel endonuclease Psc T-related clone DNA a

XX DT 03-JUN-2004 (first entry)  
 XX DE Mosaic virus movement protein gene GRAMMR clone #10.  
 XX KW ds; mismatch endonuclease; endonuclease; gene shuffling technology;  
 XX single nucleotide polymorphism; cancer susceptibility;  
 XX sequence variation redistribution; movement protein gene

XX endonuclease; molecular biology; plant propagation; phenotypic trait; herbicide tolerance; heat tolerance; cold tolerance; drought; salinity; osmotic stress; pest resistance; insect; nematode; arachnid; fungal; bacterial; viral; enzyme production; secondary metabolism; male sterility; female sterility; dwarfness; early maturity; Res I; ds. XX  
OS tobacco mosaic virus

xx  
OS Tobacco mosaic virus.  
OS Tomato mosaic virus.  
xx  
PN  
XX  
US2003157682-A1.  
yy  
PD 21-AUG-2003.

XX PN US2003148315-A1.  
XX PD 07-AUG-2003.  
XX

XX  
PR 01-FEB-2002; 2002US-0353722P  
PR 14-MAR-2002; 2002US-00098155P  
PR 01-AUG-2002; 2002US-00211079P  
XX

PF 01-AUG-2002; 2002US-00311079.  
XX  
PR 01-FEB-2002; 2002US-0353722P-  
PR 14-MAR-2002; 2002US-00098155.

PA	(PADG/)	PADGETT H. S.
PA	(VAEW/)	VAEHWONGS A. A.
PA	(VOJD/)	VOJDANI F. S.
PA	(SMIT/)	SMITH M. L.

PA (PADGETT H. S.  
PA (VAEW/) VAENHONGS A.A.  
XX  
PI  
VV  
Padgett HS, Vaewhongs AA;

PA (FITT) FITZMAURICE W. P.  
 XX Padgett HS, Vaewhongs AA, vojdani FS, Smith ML, Lindbo JA;  
 PI Fitzmaurice WP;  
 PI VV

XX  
UK  
WPI; 2003-89/548/52.  
PT  
New nucleic acid molecule encoding endonucleases, useful in molecular  
PT  
biology, specifically to generating populations of related nucleic acid  
PT  
molecules, and in plant propagation with useful phenotypic traits.

PS  
XX  
CC  
CC  
Example 15: Fig 13; 46PP; English.  
This invention relates to a novel endonuclease (Res I) nucleic acid molecule which comprises a fully defined sequence of 899 bp given in the

XX  
RS  
Example 14; SEQ ID NO 25; 79pp; English.  
XX  
CC  
The invention relates to a method of making a mismatch endonuclease

CC useful in molecular biology, and more specifically to generating CC populations of related nucleic acid molecules. They may also be used in CC plant propagation with useful phenotypic traits, such as improved CC tolerance to herbicides, improved tolerance to extremes of heat or cold, CC and increased resistance to pests.

CC bacterium with a recombinant viral vector that encodes a polynucleotide sequence for a mismatch endonuclease, growing the host so that the polynucleotide is expressed, and extracting the mismatch endonuclease enzyme from the host. The method is useful for making mismatch

CC viral; production of enzymes or secondary metabolites, male or female CC sterility, dwarfness and early maturity. The present sequence is that of CC a clone which was derived during the exemplification of the invention. XX

Query	Match	Similarity	Score	DB 10:	Length
Best Local	64.2%	Score 48.8;	DB 10;	Length 769;	
Matches 59;	77.6%	Pred. No. 4.6e-07;			
Conservative;	0;	Mismatches 0;	Indices 0;	Gaps 0	

Sequence 769 BP; 222 A; 205 C; 96 G; 246 T; 0 U; 0 other; C.C. C.C.C.C.

Best Local Similarity 77.6%; Pred. No. 4.6e-07; Matches 59; Conservative 0; Mismatches 17; Indels 0; Gaps 0; DB 12; Length 769;

Query Match 64.2%; Score 48.8; DB 12; Length 769;

Best Local Similarity 77.6%; Pred. No. 4.6e-07; Matches 59; Conservative 0; Mismatches 17; Indels 0; Gaps 0; DB 229; Score 48.8; DB 229; Length 5997;

Query Match 60.3%; Score 45.8; DB 2; Length 5997;

Best Local Similarity 76.7%; Pred. No. 8.9e-06; Matches 56; Conservative 0; Mismatches 17; Indels 0; Gaps 0; DB 229; Score 45.8; DB 229; Length 4864;

Query Match 61.1%; Score 45.8; DB 2; Length 4864;

Best Local Similarity 77.6%; Pred. No. 4.6e-07; Matches 59; Conservative 0; Mismatches 17; Indels 0; Gaps 0; DB 169; Score 48.8; DB 169; Length 154;

Query Match 61.1%; Score 45.8; DB 2; Length 154;

Best Local Similarity 77.6%; Pred. No. 4.6e-07; Matches 59; Conservative 0; Mismatches 17; Indels 0; Gaps 0; DB 169; Score 48.8; DB 169; Length 154;

RESULT 12

ADP2609/C standard; DNA; 769 BP.

XX ADP266C9;

XX ADP266C9;

XX 26-AUG-2004 (First entry)

XX DE Heteroduplex DNA #14.

XX KW Sequence variation; heteroduplex; transcription; DNA integration;

XX KW ribozyme expression; gene; ds.

XX OS Synthetic.

XX PN US2004110130-A1.

XX PD 10-JUN-2004.

XX PP 25-OCT-2002; 2002US-00230913.

XX PR 32-FEB-2001; 2001US-0265386P.

XX PR 14-FEB-2001; 2001US-0265385P.

XX PR 01-FEB-2002; 2002US-00056390.

XX PR 08-AUG-2002; 2002US-0402342P.

XX PA (LARG-) LARGE SCALE BIOLOGY CORP.

XX PI Padgett HS, Lindbo JA, Fitzmaurice WP;

XX DR WPI; 2004-440326/41.

XX PT Redistributing sequence variations between non-identical polynucleotide sequences, useful for generating improved polynucleotide having a desired characteristic, comprises making a heteroduplex and introducing a nick.

XX PS Example 15; SEQ ID NO 25; 75PP; English.

XX The invention relates to an in vitro method of redistributing sequence variations between non-identical polynucleotide sequences, comprising making a heteroduplex polynucleotide from two non-identical polynucleotides, introducing a nick in the second strand at or near a base pair mismatch site, removing the mismatched base(s) from the mismatch site where the nick occurred and using the first strand as a template to replace the removed base(s) with bases that complement the base(s) in the first strand. The invention also relates to an in vitro method of making a population of sequence variants from a heteroduplex polynucleotide sequence, obtaining a polynucleotide sequence encoding a desired functional property and identifying a reassorted DNA molecule encoding a protein with a desired functional property. The method is useful for generating an improved polynucleotide sequence or a population of improved polynucleotide sequences possessing at least one desired phenotypic characteristic (e.g., promotes transcription of linked polynucleotides), where such polynucleotides are useful for expression from a plant, animal, fungal, yeast, or bacterial expression vector, for integration to form a transgenic plant, animal or microorganism, and for expression of a ribozyme. This sequence represents DNA used in the scope of the invention.

XX Sequence 769 BP; 222 A; 205 C; 96 G; 246 T; 0 U; 0 Other;

XX SQ

RESULT 13

AAQ1218B standard; DNA; 5997 BP.

XX AAQ1218B;

XX AAQ1218B;

XX 27-AUG-2003 (revised)

XX DT 10-SEP-1991 (first entry)

XX DE Odontoglossum ring spot virus.

XX KW Odontoglossum ring spot virus; probe; coat protein; ORSV; ds.

XX OS Odontoglossum ring spot virus.

XX PN WO9102956-A.

XX PD 13-JUN-1991.

XX PP 28-NOV-1989; 89JP-00306626.

XX PR 28-NOV-1989; 89JP-00306626.

XX PA (NIOC ) NIPPON OIL KK.

XX PI Isomura K, Matsumoto Y, Chatani M, Ikegami M;

XX DR WPI; 1991-193200/26.

XX PT DNA obtld. by cleavage of cDNA corresp. to RNA - of Odontoglossum ring spot virus coding for viral coat protein and is useful as probe and vector for plant gene recombination.

XX PS Claim 2; Page 52-64; 84pp; Japanese.

XX The ds DNA is obtld. by cleavage with EcoRI of the cDNA corresp. to the genomic DNA. Organisms transformed with a vector contg. the DNA produce a peptide of mol. wt. of 33 kD, corresp. to the viral coat protein. The sequence may be used as probe, and for the prodn. of vectors for expression of plant genes. (Updated on 27-AUG-2003 to correct OS field.)

XX SQ Sequence 5997 BP; 1787 A; 1032 C; 1311 G; 1867 T; 0 U; 0 Other;

CC Query Match 60.3%; Score 45.8; DB 2; Length 5997;

CC Best Local Similarity 76.7%; Pred. No. 8.9e-06; Matches 56; Conservative 0; Mismatches 17; Indels 0; Gaps 0; DB 4805; Score 45.8; DB 4805; Length 4864;

CC Query Match 61.1%; Score 45.8; DB 2; Length 4864;

CC DB 4865; Score 45.8; DB 4865; Length 154;

CC

RESULT 14

AAQ38106 standard; cDNA to mRNA; 6597 BP.

XX

AC AAQ38106;  
 XX  
 DT 05-JUL-1993 (first entry)  
 XX  
 DE ORSV cDNA.  
 XX  
 KW Odontoglossum ring spot virus; screen; transformation; ds.  
 OS Odontoglossum ring spot virus.  
 XX  
 PN JP05030975-A.  
 XX  
 PD 09-FEB-1993.  
 XX  
 PF 26-JUL-1991; 91JP-00276075.  
 XX  
 PR 26-JUL-1991; 91JP-00276075.  
 XX  
 PA (NIOC ) NIPPON OIL KK.  
 XX  
 DR WPI; 1993-087957/11.  
 XX  
 PT cDNA of odontoglossum ring-spot virus gene - useful as vector for plants  
 and probes for screening the virus.  
 XX  
 PS Claim 2; Page 11-28; 30pp; Japanese.

XX  
 CC RNA was isolated from Odontoglossum ring spot virus and used to prepare  
 CC cDNA. The DNA or its restriction fragments can be used to screen for ORSV  
 CC or to detect genes related to ORSV. Vectors contg. the DNA sequence can  
 be used to transform E. coli, Bacillus subtilis, Arrobacterium or Plant  
 CC cells for prodn. of the recombinant 130K, 180K and 30K proteins of ORSV.  
 CC coat proteins or their peptide fragments.  
 XX  
 SQ Sequence 6597 BP; 1970 A; 1170 C; 1425 G; 2032 T; 0 U; 0 Other;  
 Query Match 50.3%; Score 45.8; DB 2; Length 6597;  
 Best Local Similarity 76.7%; Pred. No. 9.2e-06;  
 Matches 56; Conservative 0; Mismatches 17; Indels 0; Gaps 0;  
 OY 1 GTGACAGACGGCTCGCCATTGACTCTGAAAGGTGTAGGAGTCCTGGATCAA 60  
 Db 5405 GTGACAGACGGCTCGCCATTGACTCTGAAAGGTGTAGGAGTCCTGGATCAA 5464  
 OY 61 GTTACCAATGGCTG 73  
 Db 5465 GTTACCAATGGCTG 5477  
 RESULT 15  
 ID AAN30116  
 ID AAN30116 standard; RNA; 356 BP.  
 XX  
 AC AAN30116;  
 XX  
 DT 25-MAR-2003 (revised)  
 DT 02-NOV-1992 (first entry)  
 XX  
 DE TMV-RNA fragment I originating at the capped 5' end of the viral RNA and  
 DE extending into the coat protein gene.  
 XX  
 KW RNA plant virus vector; tobacco mosaic virus; ss.  
 OS Tobacco mosaic virus.  
 XX  
 FH misc\_feature 5  
 FT /\*tag= e  
 FT /label= bp No. 5400  
 FT misc\_feature 61..235  
 FT /\*tag= a  
 FT /label= nucleation region  
 FT 61..235  
 PS  
 PT RNA plant virus vector from tobacco mosaic virus etc. - for modifying  
 genes in plants to alter growth disease resistance etc.  
 XX  
 PS Example; Page 27; 56pp; English.  
 XX  
 CC The inventors claim an RNA plant virus vector from tobacco mosaic virus.  
 CC The vector comprises a nucleotide sequence originating 5' from the 5' end of  
 CC the (+) strand of the viral RNA (Fragment I) (see AAN30116; and a  
 CC sequence originating from the 3' end of the (+)strand (Fragment III) (see  
 CC AAN30114, AAN30115). The RNA vector may have foreign genetic information  
 CC inserted or attached, ultimately in the form of RNA, to the vector. The  
 CC mode of Fragment I and II production fragments enables fragments of any  
 CC desired length from any location in TMV-RNA to be generated (see  
 CC AAN30117, AAN30118). Fragment I is designated Frag. I (Pgs. The viral  
 CC replicase gene is likely included Fragment I. (Updated on 25-MAR-2003 to  
 correct PA field.)  
 XX  
 SQ Sequence 356 BP; 121 A; 48 C; 87 G; 0 T; 100 U; 0 Other;  
 Query Match 57.9%; Score 44; DB 1; Length 356;  
 Best Local Similarity 56.6%; Pred. No. 1.8e-05;  
 Matches 43; Conservative 13; Mismatches 20; Indels 0; Gaps 0;  
 OY 1 GTGACAGACGGCTCGCCATTGACTCTGAAAGGTGTAGGAGTCCTGGATCAA 60  
 OY 61 GTTACCAATGGCTG 76  
 Db 112 GUCCCUAUGUCAUCA 127  
 Search completed: December 2, 2004, 07:53:21  
 Job time : 410 secs

FT /\*tag= C  
 FT /label= coat protein gene  
 FT misc\_signal 236..320  
 FT /\*tag= b  
 FT /label= control region  
 FT misc\_feature 348..356  
 FT /\*note= "site of cleavage by ribonuclease H"  
 XX  
 PN EP67553-A.  
 XX  
 PD 22-DEC-1982.  
 XX  
 PF 27-MAY-1981; 81US-00267539.  
 XX  
 PR 27-MAY-1981; 81US-00267539.  
 XX  
 PA (CNRN ) NAT RES COUNCIL CANADA.  
 XX  
 PI Pelcher LE, Halska MC;  
 XX  
 DR WPI; 1983-00323K/01.  
 XX  
 RNA plant virus vector from tobacco mosaic virus etc. - for modifying  
 genes in plants to alter growth disease resistance etc.  
 XX  
 PS Example; Page 27; 56pp; English.  
 XX  
 CC The inventors claim an RNA plant virus vector from tobacco mosaic virus.  
 CC The vector comprises a nucleotide sequence originating 5' from the 5' end of  
 CC the (+) strand of the viral RNA (Fragment I) (see AAN30116; and a  
 CC sequence originating from the 3' end of the (+)strand (Fragment III) (see  
 CC AAN30114, AAN30115). The RNA vector may have foreign genetic information  
 CC inserted or attached, ultimately in the form of RNA, to the vector. The  
 CC mode of Fragment I and II production fragments enables fragments of any  
 CC desired length from any location in TMV-RNA to be generated (see  
 CC AAN30117, AAN30118). Fragment I is designated Frag. I (Pgs. The viral  
 CC replicase gene is likely included Fragment I. (Updated on 25-MAR-2003 to  
 correct PA field.)  
 XX  
 SQ Sequence 356 BP; 121 A; 48 C; 87 G; 0 T; 100 U; 0 Other;  
 Query Match 57.9%; Score 44; DB 1; Length 356;  
 Best Local Similarity 56.6%; Pred. No. 1.8e-05;  
 Matches 43; Conservative 13; Mismatches 20; Indels 0; Gaps 0;  
 OY 1 GTGACAGACGGCTCGCCATTGACTCTGAAAGGTGTAGGAGTCCTGGATCAA 60  
 OY 61 GTTACCAATGGCTG 76  
 Db 112 GUCCCUAUGUCAUCA 127  
 Search completed: December 2, 2004, 07:53:21  
 Job time : 410 secs

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Om nucleic - nucleic search, using sw model

Run on: December 2, 2004, 05:41:26 ; Search time 2971 Seconds

(without alignments) 932.151 Million cell updates/sec

Title: US-09-551-494-5\_COPY\_5430\_5505

Perfect score: 76

Sequence: 1 gtgacagacggctcgccaaat.....tgaagtaccaatggctgtgt 76

Scoring table: IDENTITY\_NUC

Gapov 10.0, Gapext 1.0

Searched: 3282875 seqs, 18219865908 residues

Total number of hits satisfying chosen parameters: 65645750

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-processing: Minimum Match C%

Maximum Match 100%

Listing first 45 summaries

Database : EST-\*

1: gb\_est1:\*

2: gb\_est2:\*

3: gb\_hc:\*

4: gb\_est3:\*

5: gb\_est4:\*

6: gb\_est5:\*

7: gb\_est6:\*

8: gb\_gss1:\*

9: gb\_gss2:\*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

#### SUMMARIES

Result No. Score Query Match Length DB ID Description

C 1 39.4 51.8 188 4 BM068137 KS08017C0

C 2 39.4 51.8 348 4 BM067518 KS08006E1

C 3 32.6 42.9 583 4 CB264749 41-E01466

C 4 31.6 41.6 954 8 B12288

C 5 31 40.8 389 1 AV441961

C 6 31 40.8 530 6 CR93974

C 7 31 40.8 591 9 FP020421

C 8 31 40.8 656 5 BUG6714

C 9 31 40.8 686 5 BX53578

C 10 31 40.8 1535 3 CNS0AD0

C 11 31 40.8 1579 3 CNS0ACTY

C 12 30.2 39.7 711 5 BX84538

C 13 30.2 39.7 800 5 BUG15084

C 14 30 39.5 446 9 FR0024459

C 15 29.8 39.5 561 5 BX53578

C 16 29.6 38.9 357 9 CGT742490

C 17 29.6 38.9 854 7 CO800453

C 18 29.4 38.9 732 1 AUS05571

C 19 28.8 37.9 387 5 BUG16956

C 20 28.8 37.9 398 9 CNS007V

C 21 28.8 37.9 460 7 CK09490

C 22 28.8 37.9 438 8 AZ912881

C 23 28.8 37.9 596 1 AJ763214

C 24 28.8 37.9 605 1 AJ774675

#### ALIGNMENTS

RESULT 1  
BM068137/c  
DEFINITION KS08017C0 198 bp mRNA, linear EST 11-SEP-2002  
ACCESSION BM068137  
KEYWORDS EST.  
SOURCE  
ORGANISM Capsicum annuum  
Bukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eu dicots; asterids; Lamiales; Solanaceae; Capsicum.  
1 (bases 1 to 188)  
Lee, S.-Y., Chung, Y.-H., Shin, H.-J., Goh, S.-H., Pai, H.-S., Hur, C.-G. and Choi, D.  
Generation of Expressed Sequence Tags from Hot Pepper (*Capsicum annuum* L.) and Sequence Analysis in Relation to Hypersensitive Response Against Pathogen.  
Unpublished (2001)  
COMMENT  
Contact: Doi Choi  
Genome Research Center and National Center for Genome Information  
Korea Research Institute of Bioscience and Biotechnology  
P.O. BOX 115, Yusong, Taejeon, 305-600, Republic of Korea  
Tel: 82-42-850-4240  
Fax: 82-42-850-4209  
Email: doil@mail.kribb.re.kr

FEATURES  
SOURCE  
1. .188  
/organism="Capsicum annuum"  
/mol\_type="mRNA"  
/cultivar="Hang Keun"  
/db\_xref="taxon:40722"  
/tissue\_type="anther"  
/dev\_stage="10 weeks after germination"  
/clone\_lib="X300"  
/note="Vector: pBlueScript SK(-)"

#### ORIGIN

Query Match Similarity 51.8%; Score 39.4; DB 4; Length 188;  
Best Local Similarity 71.2%; Pred. No. 0; 0; Mismatches 21; Indels 0; Gaps 0;

ORIGIN 1 GTGACGAGCTCCCAATGAACCTGAAAGCTTGAGGAGTGTGATGAA 60

Db 122 GTGTCGAGGAGGACCGTGAACCTATAGAGCAGTGTGATGAGTCATGAA 63

QY	61	GTACCAATGGCTG	73	TITLE	large-scale identification and analysis of genome-wide
Db	62		50	DEFINITION	single-nucleotide polymorphisms for mapping in <i>Arabidopsis thaliana</i>
REFERENCE		KS08000B10	KS08	JOURNAL	Genome Res. 13 (6), 1250-1257 (2003);
ACCESSION		BM067518		MEDLINE	2268320
VERSION		BM067518.1	GI:22787638	PURMED	12799357
KEYWORDS		EST.		COMMENT	Contact: Weishaar, B
SOURCE		Capsicum annuum		ADN DNA core facility at MPIZ	
ORGANISM		Capsicum annuum		Max-Planck-Institute for Plant Breeding Research	
REFERENCE		Bukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; asterids; lamids; Solanales; Solanaceae; Capsicum.		Caro-von-Linne Weg 10, 50829 Koeln, Germany	
AUTHORS	1	(bases 1 to 348)		Fax: 0049215062851	
TITLE		Lee, S., Kim, S.-Y., Chung, Y.-H., Shin, H.-J., Goh, S.-H., Pai, H.-S., Generation of Expressed Sequence Tags from Hot Pepper ( <i>Capsicum annuum</i> L.) and Sequence Analysis in Relation to Hypersensitive Response Against Pathogen		Email: weishaadempf-koeln.mpg.de	
COMMENT		Unpublished (2001)		Insert length: 583 Std Error: 0.00	
FEATURES		Contact: Doil Choi		Plate: 2 Row: B Column: 11	
source		Genome Research Center and National Center for Genome Information		Seq. primer: T7R: CTAAAGCAGCTTACATATAAGGA.	
		Korea Research Institute of Bioscience and Biotechnology		source	
		P.O. Box 115, Yusong, Taejeon, 305-600, Republic of Korea		/organism="Arabidopsis thaliana"	
		Tel: 82-42-860-4340		/mol_type="mRNA"	
		Fax: 82-42-860-4309		/db_xref="GABI:594666"	
		Email: doil@mail.kribb.re.kr		/clone="MPIZp2000B1120"	
		High quality sequence stop: 348.		/tissue_type="inflorescence"	
		Location/Qualifiers		/lab_host="E. coli TOLO"	
		1. . 348		/clone lib="MPIZ-AD10-035"	
		/organism="Capsicum annuum"		/note="Vector: PSORT; Site 1: Sali; Site 2: NotI; cDNA library from <i>Arabidopsis thaliana</i> , accession Achkarren-2; inflorescences from flower buds to young siliques; library was made at the Max-Planck-Institute for Plant Breeding Research, Cologne, Germany; cloning sites Sali-NotI, primer sites and orientation: Tr-SalI-CCCGCGRCG-Sprime-cDNA polyA-CC NotI-SP6; Note: Sequencing granted in the context of the GABI <i>Arabidopsis</i> Verbund; Genetic Diversity, establishment of high-efficiency SNP-based mapping tools and development of methods for genome-wide mutation detection; PI: Bernd Weisshaar; Sequence database: http://Gabi.rzpd.de This clone is available from RZPD; contact RZPD (clone@rzpd.de) for further information."	
ORIGIN					
RESULT	3			Query Match	42.9%; Score 32,6; DB 6; Length 383;
CB264749				Best Local Similarity	66.2%; Pred. No. 2,6; Mismatches 24; Indels 0; Gaps 0;
DEFINITION		CB264749	583 bp	Matches	47; Conservative 0; Mismatches 21; Indels 0; Gaps 0;
LOCUS		CB264749	mRNA	QY	6 AGACGGCTGCCATTGACTCACTGAAAGGTGTTGGAGGTCTGTTGATGAGTC 65
ACCESSION		41-1014660-035-002-B11-TTR	linear	Db	381 AGATGCTTCCACTGAAATCAGTGAATAGCTCTTGAAGATTTGGAGACCAATCC 440
VERSION		MP1Zp2000B1120	EST	QY	66 AATGCTGTGA 76
KEYWORDS		5-PRIME, mRNA sequence.	EST.	Db	441 AATGCTTCA 451
SOURCE		CB264749.1	Arabidopsis thaliana (thale cress)		
ORGANISM		Arabidopsis thaliana (thale cress)			
REFERENCE		Eukaryote; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.			
REFERENCE	1	(bases 1 to 583)			
AUTHORS		Schmid, K.J., Soerensen, T.R., Stracke, R., Torjek, O., Altmann, T., Mitchell-Olds, T. and Weisshaar, B.			
TITLE		Unpublished (1997)			
JOURNAL					
RESULT	4			Query Match	42.9%; Score 32,6; DB 6; Length 383;
BI12288				Best Local Similarity	66.2%; Pred. No. 2,6; Mismatches 24; Indels 0; Gaps 0;
DEFINITION		BI12288	954 bp	Matches	47; Conservative 0; Mismatches 21; Indels 0; Gaps 0;
LOCUS		BI12288	DNA	QY	6 AGACGGCTGCCATTGACTCACTGAAAGGTGTTGGAGGTCTGTTGATGAGTC 65
ACCESSION		BI12288	linear	Db	381 AGATGCTTCCACTGAAATCAGTGAATAGCTCTTGAAGATTTGGAGACCAATCC 440
VERSION		BI12288.1	EST	QY	66 AATGCTGTGA 76
KEYWORDS		GI:2093409	EST.	Db	441 AATGCTTCA 451
SOURCE			Arabidopsis thaliana (thale cress)		
ORGANISM					
REFERENCE		Eukaryote; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta; Spermatophyta; Magnoliophyta; eudicotyledons; core eudicots; rosids; eurosids II; Brassicales; Brassicaceae; Arabidopsis.			
AUTHORS	1	(bases 1 to 954)			
REFERENCE		Ecker, J., Dewar, K., Buehler, E., Kim, C., Li, Y., Shin, P., Suc, H. and			
AUTHORS		BAC End Sequences at ATGC			
TITLE		Unpublished (1997)			
JOURNAL					

COMMENT Other GSSE: T2M2-T7

COMMENT Contact: Ecker J.

Arabidopsis Thaliana Genome Center

University of Pennsylvania

Dept. of Biology, University of Pennsylvania, Philadelphia, PA 19104

Tel: 215-898-9384

Fax: 215-898-8780

Email: Jecker@genome.bio.upenn.edu

Seq primer: SP6

Class: BAC ends

High quality sequence start: 93

High quality sequence stop: 103.

FEATURES Source

1. -954 \*Qualifiers

1. .954

/organism="Arabidopsis thaliana"

/mol\_type="genomic DNA"

/ecoli\_xref="Colombia"

/db\_xref="T2M2"

/sex="hermaphrodite"

/clone="T2M2"

/clone\_lib="TAMU"

/note="Vector: pBluescriptII SK-; Site\_1: EcoRI; Site\_2: HindIII; Produced by Rod Wing"

ORIGIN

Query Match Score 41.6%; Score 31.6; DB 8; Length 954;

Best Local Similarity 56.8%; Pred. No. 6.3;

Matches 46; Conservative 0; Mismatches 25; Indels 0; Gaps 0;

Qy 6 AGAGCCTGCCAATGACTGAAAGGTTGAGGAGTCGATGAGTAC 65

Db 72 ANATGCTCTCACTGAAATCAGTAAAGCTTGAGAGATTGGAGACATTGC 131

Qy 66 AATGGCTGGA 76

Db 122 AATGGCTTAA 132

RESULT 6

CA963974 LOCUS CA963974 DEFINITION 530 bp mRNA linear EST 03-JAN-2003

DEFINITION CA963974 Infected Arabidopsis leaf Arabidopsis thaliana cDNA.

ACCESSION CA963974

VERSION CA963974.1

DEFINITION EST.

KEYWORDS EST.

SOURCE Arabidopsis thaliana (thale cress)

ORGANISM Arabidopsis thaliana

REFERENCE

AUTHORS Lundsgaard M., Emmersen J., Nielsen K.L., Wilson I., Scmerville S., and Weilinder, K.G.

TITLE EST sequencing of Erysiphe cichoracearum infected Arabidopsis

JOURNAL Unpublished (2002)

COMMENT

Contact: Karen G. Weilinder

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Sohngardsholmsvej 49, 9000 Aalborg, Denmark

TELE: +45 96358467

FAX: +45 9811808

Email: kgw@bio.auc.dk

FEATURES Source

1. -530

/organism="Arabidopsis thaliana"

/mol\_type="mRNA"

/ecoli\_xref="Colombia"

/db\_xref="T2M2"

/note="Organ: leaf; Vector: pBluescript; Mixed cDNA library of Arabidopsis and E. cichoracearum infected leaf from three weeks old Arabidopsis plants. Plants were harvested 3 days after infection and mRNA oligo dT selected."

COMMENT

Contact: Erika Asamizu

The First Laboratory for Plant Gene Research

Kazusa DNA Research Institute

Yana 153-3, Kisarazu, Chiba 292-0812, Japan

Email: asamizu@kazusa.or.jp, URL: http://www.kazusa.or.jp/en/plant/.

FEATURES Source

1. .389

/organism="Arabidopsis thaliana"

/mol\_type="mRNA"

/ecoli\_xref="Colombia"

/db\_xref="T2M2"

/db\_xref="taxon:3702"

/clone="T2M2"

/tissue\_type="aboveground organs"

ORIGIN

Query Match Score 40.8%; Score 31; DB 6; Length 530;

Best Local Similarity 64.8%; Pred. No. 9;

Matches 46; Conservative 0; Mismatches 25; Indels 0; Gaps 0;

Qy 6 AGAGGCTGCCAATGACTGAAAGGTTGAGGAGTCGATGAGTAC 65

Db 215 AGATGCTCTCACTGAAATCAGTAAAGCTTGAGAGATTGGAGACATTGC 274

Qy 66 AATGGCTGGA 76

Db 275 AATGGCTTAA 285

RESULT 7  
FR0020421  
LOCUS FR0020421  
DEFINITION *P. rubripes* GSS sequence, clone 041P11qB11, genomic survey sequence.  
ACCESSION AL013304  
VERSION AL013304.1 GI:2679672  
KEYWORDS GSS; genome survey sequence.  
SOURCE Takifugu rubripes (Fugu rubripes)  
ORGANISM Takifugu rubripes  
Bivalvia; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Acanthomorpha; Acanthopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei; Tetradontidae; Tetraodontiformes; Tetradontidae; Tetraodontidae; Takifugu.

REFERENCE 1  
AUTHORS Elgar,G., Clark,M.S., Meek,S., Smith,S., Warner,S., Edwards,Y.J., Bouchireb,N., Cottage,A., Yeo,G.S., Umrania,Y., Williams,G. and Brenner,S.

TITLE Generation and analysis of 25 Mb of genomic DNA from the pufferfish Fugu rubripes by sequence scanning  
JOURNAL Genome Res. 9 (10), 960-971 (1999)  
MEDLINE 9945097  
PUBMED 1052524  
REFERENCE 2  
AUTHORS Elgar,G., Clark,M., Smith,S., Meek,S., Warner,S., Umrania,Y., Williams,G. and Brenner,S.  
TITLE Direct Submission  
JOURNAL Submitted (08-DEC-1997) MRC Human Genome Mapping Project Resource Centre Hinxton, Cambridge, CB10 1SB. Email: biohelp@hgmpr.mrc.ac.uk  
COMMENT Vector: pBluescript II KS  
PRIMER: phagemid  
DESCR: KS

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Db 423 ATTCCTTTA 433

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ACCESSION BU636451  
VERSION BU636451.1 GI:23303706  
KEYWORDS EST.  
SOURCE Arabidopsis thaliana (thale cress)

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Db 113 GGGCTGAGTGTGACTCTGGANATGTTGACGATCTGGGAGACGTGTAAG 172  
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Db 173 GTGTCA 179

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REFERENCE Lundsgaard,M., Brummersen,J., Nielsen,K.L., Wilson,I., Somerville,S.  
AUTHORS and Welinder,K.G.  
TITLE EST sequencing of *Erysiphe cichoracearum* infected *Arabidopsis* plants  
JOURNAL Unpublished (2002)  
COMMENT Contact: Karen G. Welinder  
Institut for bioteknologi  
Aalborg Universitet  
Søgaardstrømsvej 49, 9000 Aalborg, Denmark  
Tel: +45 96358467  
Fax: +45 98141808  
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AUTHORS Lundsgaard,M., Brummersen,J., Nielsen,K.L., Wilson,I., Somerville,S.  
REFERENCE Lundsgaard,M., Brummersen,J., Nielsen,K.L., Wilson,I., Somerville,S.  
AUTHORS and Welinder,K.G.

library of *Arabidopsis* and *B. cichoracearum* infected leaf tissue from three weeks old *Arabidopsis* plants. Plants were harvested 3 days after infection and mRNA oligo dt selected.

GenCore version 5.1.6  
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Om nucleic - nucleic search, using sw model

Run on:

December 1, 2004, 20:23:09 ; Search time 1635 Seconds  
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2198.176 Million cell updates/sec

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Gapop 10.0 , Gapext 1.0

Searched:

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Total number of hits satisfying chosen parameters: 9053458

Minimum DB seq length: 0

Maximum DB seq length: 200000000

Post-Processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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8: gb\_pcl:\*

9: gb\_pcr:\*

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13: gb\_uni:\*

14: gb\_vrl:\*

### SUMMARIES

Pred. No is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

### ALIGNMENTS

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 REFERENCE  
 AUTHORS Metzlaaff, M.H., Goosselle, V.M., Meulewaeter, F. and Fache, J.C.  
 TITLE Improved methods and means for delivering inhibitory rna to plants and applications thereof  
 JOURNAL Patent: WO 03052108-A 7 26-JUN-2003;  
 FILER Bayer BioScience N.V. (BE)  
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 AUTHORS Garcia-Arenal, F.  
 TITLE Sequence and structure at the genome 3' end of the U2-strain of tobacco mosaic virus, a histidine-accepting tobamovirus  
 JOURNAL Virology 167 (1), 201-206 (1988)  
 MEDLINE 8945644  
 PUBLISHED 3-8-1986  
 REFERENCE 2 (bases 1 to 6355)  
 AUTHORS Solis, I. and Garcia-Arenal, F.  
 TITLE The complete nucleotide sequence of the genomic RNA of the tobacco mosaic tobacco mild green mosaic virus  
 JOURNAL Virology 177 (2), 553-558 (1991)  
 MEDLINE 9032027  
 PUBLISHED 23-7-1991  
 COMMENT  
 Original source text: Tobacco mild green mosaic virus (strain U2-TMV), cDNA, to viral RNA, from Nicotiana tabacum cv. Samsum. Draft entry and computer-readable sequence for [1] kindly submitted by F. Garcia-Arenal, 10-FEB-1989. The RNA appears to have a tRNA-like, L-shaped structure at the 3' terminus, linked to a quasi-continuous double-helical stalk, with five pseudoknots involved in the formation of the whole structure. However, the structure of U2-tRNA is less stringently conserved than the 3' termini of ,vulgaris, and other histidine-accepting tobamoviruses. Draft entry and computer-readable sequence for [1] kindly submitted by F. Garcia-Arenal, 08-MAY-1990, for release after publication.  
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## ORIGIN

Query Match Similarity 60.3%; Score 45.8; DB 6; Length 5997;  
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Db 613 GTAACTGAGAGGCCACCGAACCTACTGAAAGGTGTTGAGGAGTCGAGAA 672  
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Db 673 GTTCCTATGGCTG 685

## RESULT 8

LOCUS E03624 6597 bp RNA linear PAT 29-SEP-1997  
DEFINITION DNA encoding Odontoglossum ring spot virus (ORSV) genomic RNA.  
ACCESSION E03624  
VERSION E03624.1 GI:2172508  
KEYWORDS  
SOURCE  
ORGANISM Odontoglossum ringspot virus  
REFERENCE 1 (bases 1 to 6597)  
AUTHORS Isomura, Y., Matsumoto, Y., Chatani, M., Mizuta, Y. and Ikegami, M.  
TITLE CDNA OF ORSV GENE  
JOURNAL Patent: JP 199303975-A 1 09-FEB-1993;  
NIPPON OIL CO LTD  
OS Odontoglossum ring spot virus  
PN JP 199303975-A/1  
PD 09-FEB-1993  
PF 26-JUL-1991 JP 1991276075  
PI ISOMURA, YOSHIKATSU, MATSUMOTO YOSHITOMO, CHATANI MASAHI, PI  
MIZUTA, YOSHIMORI, IKEGAMI MASAHI  
PC C12N15/40, C07K15/04, C12N1/21, C12N15/11, C12N15/70, C12P21/02, PC  
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## RESULT 9

LOCUS E04305 6597 bp RNA linear PAT 29-SEP-1997  
DEFINITION DNA encoding Odontoglossum ring spot virus (ORSV) genomic RNA.  
ACCESSION E04305  
VERSION E04305.1 GI:2172508  
KEYWORDS  
SOURCE  
ORGANISM Odontoglossum ringspot virus  
REFERENCE 1 (bases 1 to 6597)  
AUTHORS Isomura, Y., Matsumoto, Y., Chatani, M., Mizuta, Y. and Ikegami, M.  
TITLE CDNA OF ORSV GENE  
JOURNAL Patent: JP 199303975-A 1 09-FEB-1993;  
NIPPON OIL CO LTD  
OS Odontoglossum ring spot virus  
PN JP 199303975-A/1  
PD 09-FEB-1993  
PF 26-JUL-1991 JP 1991276075  
PI ISOMURA, YOSHIKATSU, MATSUMOTO YOSHITOMO, CHATANI MASAHI, PI  
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CC hypothetical: No;  
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LOCUS	AY571290					
DEFINITION	Odontoglossum ringspot virus strain Taiwan, complete genome.					
ACCESSION	AY571290					
VERSION	AY571290.1					
KEYWORDS	.					
SOURCE	Odontoglossum ringspot virus					
ORGANISM	Virus, ssRNA, positive-strand viruses, no DNA stage; Tobamovirus.					
REFERENCE	1. (bases 1 to 6612)					
AUTHORS	Wang, H. L. and Wang, J. N.					
TITLE	Molecular sequencing and analysis of the viral genome of					
JOURNAL	Odontoglossum ringspot virus Taiwan strain					
REFERENCE	Zhi Wu Bing Li Xue Hui Kan 13 (2004) In press					
AUTHORS	2. (bases 1 to 6612)					
TITLE	Wang, H. L. and Wang, J. N.					
JOURNAL	Direct Submission					
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Query Match 57.9%; Score 44; DB 14; Length 6395;  
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